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by

Jack Y. Peng, Mark R. van den Bergh and William C. Harrigan, Jr.

on the invention entitled

THREE-PHASE NANOCOMPOSITE

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prepared by:

ROZSA & CHEN LLP

Attorneys At Law

15910 Ventura Boulevard, Suite 1601

Encino, California 91436-2815

Telephone (818) 783-0990

Facsimile (818) 783-0992

e-mail: counsel@rozsa-chen.com

THREE-PHASE NANOCOMPOSITE

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention generally relates to the field of industrial production of aluminum alloys, and more particularly the field of industrial production of aluminum metal matrix nanocomposites

2. Description of the Prior Art

Currently 2000, 4000 and 6000 series aluminum alloys are widely used for lightweight structural applications that do not require high material strength and high specific stiffness or modulus. Highly loaded structural applications produced from aluminum alloys typically use 7000 series alloys. Aluminum alloys in the 7000 series are also more difficult to form into final products than competing 2000 and 6000 series aluminum alloys.

Particularly, Micro Ceramic Particulate Reinforced Aluminum Matrix Composites (MCP-AMCs) have been used in the aerospace industry and other areas due to their high specific stiffness. Aluminum oxide, boron carbide and silicon carbide micro size powders are the most common reinforcements. The specific stiffness increase is dictated by the properties of the ceramic used. Boron carbide has the highest elastic modulus and the lowest density of these reinforcements.

1 Silicon carbide, which has a slightly lower elastic modulus and higher
2 density than boron carbide, yields a MCP-AMC with intermediate properties.
3 Aluminum oxide has the highest density and the lowest modulus among the three
4 ceramics. Therefore, Al_2O_3 reinforce aluminum possesses the lowest specific
5 properties .
6

7 The strength of a MCP-AMC is mainly determined by the strength of the
8 aluminum matrix alloy. The strength of a MCP-AMC using a 6000 series
9 aluminum alloy matrix is not as high as the strength of 7000 series aluminum. For
10 example, 15 volume % SiC reinforced 6061 aluminum has a yield strength of 58
11 ksi., which is lower than the 62 ksi. for 7075-T76 aluminum alloy.
12

13 It has been shown that Nano Ceramic Particulate Reinforced Aluminum
14 Matrix Composites (NCP-AMCs) have higher strength than that of MCP-AMCs.
15 Uniformly distributed nano-scale ceramic particles in an aluminum matrix enhance
16 the strength of the resultant composite. For example, a uniform distribution of
17 about 0.5 volume % of nano-scale aluminum oxide in 6092 aluminum increases the
18 strength of the NCP-AMC about 10%. The same amount of micro ceramic powder
19 uniformly distributed in a pure aluminum alloy has no effect to the strength.
20

21 United States Patent Application Serial No. 10/738,275 by Peng *et al.*
22 disclosed how to produce nano aluminum oxide reinforced aluminum matrix
23 composites, which can have up to about 45 vol. % nano phase Al_2O_3 reinforcement.
24 However, the specific stiffness or specific modulus of such nano aluminum
25 composite is lower than that of a micro B_4C or SiC powder reinforced aluminum
26 matrix composite.
27
28

SUMMARY OF THE INVENTION

The present invention is a novel and unique three-phase nanocomposite.

The primary object of the present invention is to create a novel and unique nanocomposite system that has high strength and high specific stiffness.

Described briefly, the present invention is a three-phase nanocomposite that comprises about 0.5 to 10 vol % nano-scale aluminum oxide particles and about 1 to 45% high modulus ceramic particles and an aluminum alloy matrix. The nano phase is to enhance nanocomposite strength and the modulus phase is to enhance the specific modulus of the resulting nanocomposite.

Further novel features and other objects of the present invention will become apparent from the following detailed description, discussion and the appended claims.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Although specific embodiments of the present invention will now be described, it should be understood that such embodiments are by way of example only and merely illustrative of but a small number of the many possible specific embodiments which can represent applications of the principles of the present invention. Various changes and modifications obvious to one skilled in the art to which the present invention pertains are deemed to be within the spirit, scope and contemplation of the present invention as further defined in the appended claims.

The nanocomposite of this invention is a three-phase nanocomposite system. The first phase is an aluminum alloy matrix phase. The second is a nano aluminum oxide phase that provides strength enhancement. The third is a micro-scale ceramic particle phase possessing a higher modulus than aluminum oxide that contributes to stiffness enhancement. For simplification, the aluminum alloy matrix phase is called the matrix phase, the nano aluminum oxide phase is called the nano phase and the high modulus ceramic phase is called the modulus phase. The nano phase and the modulus phases are uniformly distributed throughout the matrix to form the nanocomposite that has high strength and high specific stiffness.

The matrix phase can be 2000, 6000 or 7000 series aluminum as defined by the Aluminum Association, or other aluminum alloys. The modulus phase can be boron carbide, silicon carbide or other ceramic powders that have higher modulus than aluminum oxide. The modulus phase can be in the nano-scale or micro-scale range. The nano-scale modulus phase does not offer better specific stiffness enhancement than the micro-scale modulus phase. The micro-scale phase is also preferred for better producibility.

1 The average particle size of the micro-scale modulus phase is from about 0.2
2 to 15 microns. The volume loading of the nano phase is about 0.5 to 10 percent.
3 The modulus phase in the composite has about 1 to 45 volume percent. Examples
4 of the material system compositions for the three-phase nanocomposite of this
5 invention are contained in Table 1.

6
7 The three-phase nanocomposite can be produced by consolidating a uniform
8 mixture of aluminum alloy powder, nano aluminum oxide powder and the micro-
9 scale ceramic powder.

10
11 A low-cost manufacturing method of producing the three-phase
12 nanocomposite is to modify the manufacturing method of nanocomposites
13 disclosed in United States Patent Application Serial No. 10/738,275 by Peng *et al.*
14 The major sequential steps of the manufacturing method of this invention are
15 described as the following:

16
17 **Step 1. Preparing the Nano- Al_2O_3 -surface Aluminum Powder for the**
18 **Nano Phase**

19
20 The nano phase is created by breaking the nano aluminum oxide layer on the
21 aluminum powder surface into a nano-scale reinforcement for strength
22 enhancement. The total volume percent of the Al_2O_3 nano-scale phase, N_T , and the
23 total volume percent of the modulus phase, V_M , in a three-phase nanocomposite are
24 predetermined according to the strength and modulus requirement to be met as well
25 as the aluminum matrix alloy type. Different applications may require different N_T
26 and V_M . N_T is about 0.5 to 10 vol % and V_M is about 1 to 45 vol %. The volume
27 percent of nano phase Al_2O_3 , N_{Al} , needed in the aluminum powder can be
28 calculated by Equation [1]:

TABLE 1
THREE-PHASE NANOCOMPOSITE SYSTEM

Three-Phase Nanocomposite

Matrix Phase		Nano Phase			Modulus Phase		
Metal	Chemicals	Ceramic	Particle size (nm)	Volume %	Ceramic	Particle size (m)	Volume %
Aluminum alloy	2000, 6000, and 7000 series or other Al alloys	Al ₂ O ₃	~10 – 800	~ 0.5 - 10	B ₄ C, SiC or other high modulus ceramics	~0.2 – 15	~1 - 45

$$N_{Al} = N_T(1 + V_M/V_{Al}) \quad [1]$$

The total volume percentage of nano-scale Al_2O_3 on a spherical shaped aluminum powder, the size of average aluminum powder particle size D and the thickness of the nano -scale Al_2O_3 layer have a relationship defined by Equation [2]:

$$N_{Al} = 1 - (1 - 2T/D)^3 \quad [2]$$

For a given thickness of the nano -scale Al_2O_3 surface layer, the smaller average particle size, the more total Al_2O_3 is formed.

The volume percent of Al_2O_3 in an aluminum powder of irregular shape is also a function of the average particle size of the powder and the thicknesses of Al_2O_3 layer. The relationship between Al_2O_3 volume percentage, particle size and layer thickness for irregularly shaped aluminum powder will be similar to that found in spherical particles. A geometric factor is required to account for the greater surface to volume for the nonspherical particles.

Spherical shaped aluminum powder is used in this invention because it is easier to analyze. A spherical shape has the smallest surface to volume ratio, which provides the highest thickness of Al_2O_3 for a given particle size and total oxide content.

This invention uses the Equation [1] to determine the volume percentage of nano-scale Al_2O_3 layers needed for a three-phase nanocomposite. Equation [2] is used to control the production of aluminum powder with a nano-scale Al_2O_3 surface layer.

1 There are generally three ways to control the Al_2O_3 volume percent in the
2 aluminum powder.

3
4 (1) For existing aluminum powder with a fixed thickness Al_2O_3 surface
5 layer:

- 6
7 (a) Determining the Al_2O_3 layer thickness of aluminum powder
8 produced by an aluminum powder manufacturer. The thickness
9 can be indirectly measured. First, measure the total Al_2O_3
10 percentage of the aluminum with a known average particle size.
11 Calculate the thickness of the Al_2O_3 layer by Equation [2];
12 (b) Using Equation [2] again, calculate the average particle size
13 needed for a predetermined Al_2O_3 vol %; and
14 (c) Classify the aluminum powder to the same average particle size
15 as the calculated value.

16
17 (2) For existing aluminum powder that possesses a large average particle
18 size:

- 19
20 (a) Measuring the average particle size;
21 (b) Determining the existing Al_2O_3 layer thickness by the method
22 in (1)(a);
23 (c) Determining the required thickness of Al_2O_3 layer by the
24 Equation [2] for a predetermined Al_2O_3 vol %; and
25 (d) Increasing the Al_2O_3 layer thickness if the existing thickness is
26 less than the required thickness. The simplest way is to heat
27 treat the aluminum powder in dry air at a temperature below the
28 aluminum's melting point.

1 A fluidized bed furnace is an efficient method for increasing the thickness of
2 the Al_2O_3 layer. The Al_2O_3 thickness is controlled by the temperature and the
3 furnace cycle time.

4
5 **Step 2. Preparing Powder Mixture**

6
7 The second step is to create a powder mixture that contains all necessary
8 elements required to create the three-phase nanocomposite.

9
10 The modulus phase powder is selected from boron carbide, silicon carbide
11 and other ceramic powders that have a modulus higher than aluminum oxide. The
12 average particle size of the modulus phase ceramic powder is from about 0.2 to 15
13 microns with a normal particle size distribution. The chemical purity of the
14 modulus phase ceramic powder is greater than about 98%.

15
16 The nano phase will be created by breaking the nano Al_2O_3 surface layers on
17 the aluminum particles during the consolidation step (Step 3) and subsequent
18 metalworking.

19
20 The matrix phase is the aluminum alloy. The aluminum alloy is selected
21 from aluminum alloys containing elements taken from aluminum, boron, cobalt,
22 copper, iron, magnesium, manganese, nickel, silicon, titanium, zinc, alloys and
23 combination thereof. There are two routes to prepare the aluminum alloy matrix:

24
25 (1) Aluminum alloy powder with nano-scale surface aluminum oxide:

26
27 By following the procedure in Step 1, the aluminum alloy powder with
28 a predetermined percentage of nano-scale surface aluminum oxide can be produced

1 by atomizing a desired aluminum alloy ingot. Such aluminum alloy powder is
2 ready to be mixed with the modulus phase ceramic powder.

3
4 (2) Mixing alloy elemental powders with an unalloyed aluminum powder
5 with a nano-scale surface oxide:
6

7 The powder metallurgy method of creating an aluminum alloy by blending
8 elemental powders is employed when unalloyed aluminum powder with a nano-
9 scale surface aluminum oxide is used. The unalloyed aluminum powder, alloy
10 elemental powders and the ceramic powder are prepared in Step 2(1) and the
11 constituents are uniformly mixed using a blender, jet mill or ball mill to form a
12 powder mixture.
13

14 The alloy elemental powders are selected from the group consisting of
15 boron, cobalt, copper, iron, magnesium, manganese, nickel, silicon, titanium, zinc,
16 alloys and combination thereof. It is preferred that the average particle sizes of the
17 alloy elemental powders be in the same particle size range as that of the unalloyed
18 aluminum powder.
19

20 The Al_2O_3 volume percent should be adjusted according to the percentage of
21 additional alloy elemental powder being used.
22

23 **Step 3. Consolidating the Powder Mixture**
24

25 The powder mixture is loaded into a billet tool and is pressed at room
26 temperature to form a compacted mixture that has between about 50% and 95% of
27 theoretical density. The billet tool size can be very small or sized for compacting
28 quantities approaching about 1000 kg (2,200 lb) of powder.

1 There are various powder metallurgy methods that can be employed to
2 consolidate the compacted mixture. The following are typical processes:

3
4 (1) Vacuum/Inert-gas/Air Hot-Pressing:

5
6 Under vacuum, inert-gas or air, the compacted mixture is heated to a
7 degassing temperature range and then is held at this range for more than about one-
8 half hour for degassing purposes. The degassing temperature range is from
9 between about 230°C (450°F) and less than the lowest eutectic melt temperature of
10 any elemental alloy constituent powder in the matrix metal. The main function of
11 degassing is to remove H₂O from the metal powder mixture.

12
13 After the degassing period, the temperature is raised to the consolidation
14 temperature. The highest consolidation temperature is the highest eutectic melt
15 temperature of any alloy constituent powder in the metal matrix. However, the
16 actual consolidation temperature may be below this eutectic melt temperature.
17 The consolidation temperature is lower than the aluminum melting point.
18 The consolidation temperature is typically between about 230 and 615°C (450 and
19 1145°F). While the consolidation temperature and vacuum, inert-gas or air
20 environment is maintained, the degassed, compacted mixture is pressed to full
21 density resulting in a composite billet. In the case of a thin Al₂O₃ layer,
22 consolidation of the powder should take place under vacuum or inert gas to prevent
23 additional oxidation.

24
25 (2) Cold Isostatic Press/Sinter:

26
27 In the Cold Isostatic Press (CIP)/Sinter process, the powder mixture is
28 compacted at room temperature in Step 3 to about 85% to 95% of theoretical

1 density. Pressing the powder mixture to high density at room temperature requires
2 pressures between about 50,000 psi and 85,000 psi. Typically, a Cold Isostatic
3 Press is employed.

4
5 The compacted mixture is then sintered in vacuum, in inert-gas, or in air.
6 The compacted mixture is heated to the degassing temperature range and then is
7 held at this temperature range for more than about one-half hour to be degassed.
8 After degassing, the degassed compacted mixture is heated to a sintering
9 temperature that is the highest eutectic melt temperature of the elemental powder in
10 the matrix metal so that sintering of the matrix takes place to form the composite
11 billet. This sintered composite billet has a density that is still approximately that of
12 the starting compacted mixture, between 85% and 95% of the theoretical density,
13 but is sealed by the sintering process. This sealing is needed to avoid internal
14 oxidation of alloy elements in the billet during subsequent hot working. The
15 composite will have 98% to 100% of the theoretical density after following metal
16 working.

17
18 In the case of aluminum powders with a thin Al_2O_3 layer, sintering should
19 take place under vacuum or inert gas to prevent additional oxidation.

20
21 (3) Cold Compacting/Hot Press:
22

23 In the case of alloyed aluminum powder with a thick Al_2O_3 layer, the powder
24 mixture can be consolidated without degassing since the thick Al_2O_3 layer prevents
25 additional oxidization that is detrimental to the resultant composite. The mixture is
26 loaded into a billet tool and is pressed at room temperature to form a compacted
27 mixture that is between about 50% and 95% of the theoretical density.
28

1 The compacted mixture is heated to the consolidation temperature in
2 vacuum, inert-gas or air. While the consolidation temperature and vacuum, inert-
3 gas, or air continues to be maintained, the compacted mixture is hot pressed to
4 approximately 98 to 100% of theoretical density to produce the composite billet.

5
6 Boron carbide and silicon carbide are harder than aluminum oxide. B_4C and
7 SiC powder particles have irregular shapes and sharp edges. There is great shear
8 and crash forces between aluminum powder and modulus ceramic powder to help
9 break the Al_2O_3 layers on surfaces of aluminum powder into nano Al_2O_3 particles.

10
11 **Step 4. Metal Working the Composite Billet**

12
13 Most of the Al_2O_3 surface layers are broken up during billet consolidation
14 due to their brittle nature. Unaffected Al_2O_3 surface layers are broken up during
15 cold or hot metal working, or any combination of these steps.

16
17 The cold work can be cold extrusion, cold forging, cold rolling or any
18 combination of these steps. During hot metal working, the composite billet is
19 plastically deformed at a temperature between about 230 and 615°C (450 and
20 1145°F), which is below the melting point of aluminum. The hot metal working
21 can be hot extrusion, hot forging or hot rolling or any combination of these steps.

22
23 The metal working can be incorporated as part of fabrication processes for
24 producing final three-phase nanocomposite products. After the consolidation and
25 metal working, the Al_2O_3 surface layers of nano-scale thickness are broken into
26 a nano-scale Al_2O_3 particulate reinforcement phase which is uniformly distributed
27 in the composite resulting in a homogenous three-phase nanocomposite.

1 Described generally, the present invention is a nanocomposite comprises
2 an aluminum alloy phase, a nano-scale aluminum oxide phase, and a modulus
3 phase.

4
5 Particularly, in the present invention three phase nanocomposite described
6 above: (1) the aluminum alloy includes elements taken from aluminum, boron,
7 cobalt, copper, iron, magnesium, manganese, nickel, silicon, titanium, zinc, alloys
8 and combination thereof; (2) the nano-scale aluminum oxide phase is nano-scale
9 aluminum oxide particles having an average particle size between about 10 nm to
10 about 800 nm and being uniformly distributed in the nanocomposite; and (3) the
11 modulus phase is ceramic particles selected from boron carbide powder, silicon
12 carbide powder or other ceramic powders having higher elastic modulus than that
13 of aluminum oxide and having an average particle size between about 0.2 microns
14 to about 15 microns and being uniformly distributed in the nanocomposite.

15
16 Preferably, the present invention nanocomposite described above has about
17 0.5 to about 10 volume percentage of the nano aluminum oxide particles, and about
18 1 to about 45 volume percentage of the modulus ceramic particles.

19
20 Of course the present invention is not intended to be restricted to any
21 particular form or arrangement, or any specific embodiment, or any specific use,
22 disclosed herein, since the same may be modified in various particulars or relations
23 without departing from the spirit or scope of the claimed invention hereinabove
24 shown and described of which the apparatus or method shown is intended only for
25 illustration and disclosure of an operative embodiment and not to show all of the
26 various forms or modifications in which this invention might be embodied or
27 operated.

28

1 The present invention has been described in considerable detail in order to
2 comply with the patent laws by providing full public disclosure of at least one of
3 its forms. However, such detailed description is not intended in any way to limit
4 the broad features or principles of the present invention, or the scope of the patent
5 to be granted. Therefore, the invention is to be limited only by the scope of the
6 appended claims.

7

8 **WHAT IS CLAIMED IS:**

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